
REMARKS

This paper is filed concurrently with the application papers requesting filing and immediate examination of a United States national application based on international application no. PCT/US2004/000892, January 13, 2004, claiming priority to US provisional application 60/440,812 filed January 17, 2003. Please enter the foregoing amendments prior to calculation of the filing fee and prior to examination on the merits.

The first paragraph of the specification has been amended herein to provide updated information regarding cross-reference to related applications under 37 C.F.R. § 1.78.

Applicant has amended Claim 1 (positions 5, 8, and 14) and paragraphs 24, 27, and 33 of the specification to delete the initial reference to "Inc" (indoline-2-carboxylic acid) and include it within the listing of D- or L- amino acids. Support for the amendment is apparent from the original claim and paragraphs as well as in original Claim 1 and paragraphs 24, 27, and 33 of the priority document, US provisional application 60/440,812.

Applicant has amended Claim 1 (position 22) and paragraph 41 of the specification to include the amino acid alanine (Ala). Support for the amendment can be found on page 13, paragraph 132 of the international publication WO 2004/066966 as well as paragraph 131 of the priority document, US provisional application 60/440,812.

Applicant has amended Claim 1 (position 23) and paragraph 42 of the specification to include the amino acid tryptophan (Trp). Support for the amendment can be found on page 13, paragraph 133 of the international publication WO 2004/066966 as well as paragraph 132 of the priority document, US provisional application 60/440,812.

Applicant has amended Claim 1 (position 25) and paragraph 44 of the specification to delete the duplicate reference to α -aminoisobutyric acid (Aib) and insert therefor substituted amino acid "Apc". Support for the amendment can be found on page 12, paragraph 99 of the international publication WO 2004/066966 as well as in Claim 1 and paragraph 43 of the priority document, US provisional application 60/440,812.

Applicant has amended Claim 1 and paragraph 66 of the specification to insert amino acid position "A³⁴" between A³³ and A³⁵. Support for the amendment can be found on page 9, paragraph 53 of the international publication WO 2004/066966 as well as in Claim 1 and paragraph 52 of the priority document, US provisional application 60/440,812.

Applicant has amended Claims 9 and 12, as well as paragraphs 393 and 426 of the specification to include the sequence Ac-(A6c²⁴)hPYY(24-36)NH₂ (SEQ ID NO. 64). Support for the

amendment can be found throughout the application, see, e.g., paragraph 305 of the international publication WO 2004/066966.

Last, Applicant requests amendment to the table found on page 37 of the application. After filing, Applicant noticed that the purity data from compounds 26, 27, and 28 had inadvertently been switched: the purity value set forth for compound 26 (97.5) was actually the reading for compound 27; the purity value set forth for compound 27 (96.3) was actually the reading for compound 28; and the purity value set forth for compound 28 (98.8) was actually the reading for compound 26.

The remaining amendments to the claims are clerical in nature, e.g., they either eliminate improper multiple dependencies or clarify the claim language to aid examination without substantively changing the claim. The claims are thus presented in proper form for review in the USPTO. It is not the intention of the Applicants to abandon any of the inventive subject matter disclosed in the application as originally filed. Support for these amendments is apparent in the original claims 1-28, and no new matter is added thereby.

A check (No. 6462) in the amount of \$1,000.00 to cover the national stage filing fees accompanies these papers, and the fees are calculated taking the foregoing amendments into account, since the amendments eliminate some multiple dependencies from the claims. No additional fees are believed to be due; however, the Commissioner is specifically authorized to charge any additional fees deemed to be necessary in connection with the filing of this paper or any of the accompanying papers, or to charge other fees necessary to complete this US filing to Deposit Account 50-0268.

Entry of the foregoing amendments and examination of the claims as amended are requested.

Respectfully submitted,



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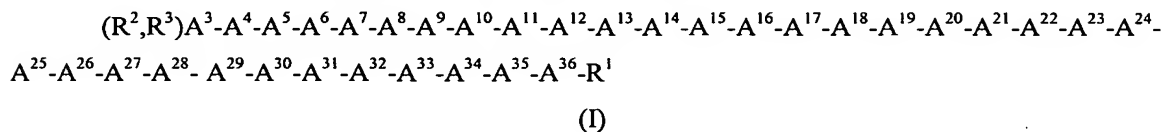
IN THE CLAIMS

COMPLETE LISTING OF ALL CLAIMS, WITH MARKINGS AND STATUS IDENTIFIERS
 (Currently amended claims showing deletions by ~~striketrough~~ and additions by underlining)

This listing of claims will replace all prior versions and listings of the claims in the application.

Listing of Claims:

1. (currently amended) A compound according to formula (I):



wherein:

A³ is Acc, Act, or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Ile, Leu, Nle, Tle, hLeu, Cha, Val, Ala, Nva, and Abu, or the N-methylated variant of Acc, Act, or Aib, or of said D- or L- amino acid, or is deleted;

A⁴ is Aib, Acc, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Lys, Arg, hArg, Orn, Dab, Dap, and HN-CH((CH₂)_n-N(R⁴R⁵))-C(O), or the N-methylated variant of Aib, Acc, or Apc, or of said D- or L- amino acid, or is deleted;

A⁵ is ~~Inc~~, or a D- or L- amino acid selected from the list of amino acids consisting of Pro, Thz, Dmt, Dhp, Ktp, 4Hyp, 3Hyp, Pip, Tic, Inc and Oic, or the N-methylated variant of Inc or of said D- or L- amino acid, or is deleted;

A⁶ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Glu, Asp, Gln, Asn, Lys, Arg, Orn, Dab, Dap, and hArg, or the N-methylated variant of Acc or Aib, or of said D- or L- amino acid, or is deleted;

A⁷ is Acc, Act, Aib, Apc, or Gly, or a D- or L- amino acid selected from the list of amino acids consisting of Ala, Abu, Val, and Nva, or the N-methylated variant of Acc, Act, Aib, Apc, or Gly, or of said D- or L- amino acid, or is deleted;

A⁸ is ~~Inc~~, or a D- or L- amino acid selected from the list of amino acids consisting of Pro, Thz, Dmt, Dhp, Ktp, 4Hyp, 3Hyp, Pip, Tic, Inc and Oic, or the N-methylated variant of Inc or of said D- or L- amino acid, or is deleted;

A⁹ is Acc, Aib, or Gly, or D- or L- Ala, or the N-methylated variant of Acc, Aib, Gly, or D- or L- Ala, or is deleted;

A¹⁰ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Glu, Asp, Gln, and Asn, or the N-methylated variant of Acc or Aib, or of said D- or L- amino acid, or is deleted;

A¹¹ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Asp, Glu, Gln, and Asn, or the N-methylated variant of Acc or Aib, or of said D- or L- amino acid, or is deleted;

A¹² is Acc, Act, Aib, Apc, or Gly, or a D- or L- amino acid selected from the list of amino acids consisting of Ala, Abu, Val, and Nva, or the N-methylated variant of Acc, Act, Aib, Apc, or Gly, or of said D- or L- amino acid, or is deleted;

A¹³ is Acc, Aib, or Act, or a D- or L- amino acid selected from the list of amino acids consisting of Ser, Thr, Ala, Abu, and Val, or the N-methylated variant of Acc, Aib, or Act, or of said D- or L- amino acid, or is deleted;

A¹⁴ is ~~Inc or~~ a D- or L- amino acid selected from the list of amino acids consisting of Pro, Thz, Dmt, Dhp, Ktp, 4Hyp, 3Hyp, Pip, Tic, Inc and Oic, or the N-methylated variant of Inc or of said D- or L- amino acid, or is deleted;

A¹⁵ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Glu, Asp, Gln, and Asn, or the N-methylated variant of Acc or Aib, or of said D- or L- amino acid, or is deleted;

A¹⁶ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Glu, Asp, Gln, and Asn, or the N-methylated variant of Acc or Aib, or of said D- or L- amino acid, or is deleted;

A¹⁷ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Leu, Ile, Nle, Tle, hLeu, Cha, Val, Ala, Nva, Abu, and Phe, or the N-methylated variant of Acc or Aib, or of said D- or L- amino acid, or is deleted;

A¹⁸ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Asn, Gln, Glu, and Asp, or the N-methylated variant of Acc or Aib, or of said D- or L- amino acid, or is deleted;

A¹⁹ is Acc, Aib, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Arg, hArg, Lys, Orn, Dab, Dap, and $\text{HN-CH}((\text{CH}_2)_n\text{-N(R}^4\text{R}^5))\text{-C(O)}$, or the N-methylated variant of Acc, Aib, or Apc, or of said D- or L- amino acid, or is deleted;

A²⁰ is Acc or Aic, or a D- or L- amino acid selected from the list of amino acids consisting of Tyr, Phe, hPhe, 2Thi, 3Thi, Taz, 2Fua, Trp, 2Nal, 1Nal, Cha, 2Pal, 3Pal, 4Pal, and $(\text{X}^1, \text{X}^2, \text{X}^3, \text{X}^4, \text{X}^5)\text{Phe}$, or the N-methylated variant of Acc or Aic, or of said D- or L- amino acid, or is deleted;

A²¹ is Acc or Aic, or a D- or L- amino acid selected from the list of amino acids consisting of Tyr, Phe, hPhe, 2Thi, 3Thi, Taz, 2Fua, Trp, 2Nal, 1Nal, Cha, 2Pal, 3Pal, 4Pal, and (X¹,X²,X³,X⁴,X⁵)Phe, or the N-methylated variant of Acc or Aic, or of said D- or L- amino acid, or is deleted;

A²² is Acc, Act, Aib, Apc, or Gly, or a D- or L- amino acid selected from the list of amino acids consisting of Ala, Aib, Abu, Val, and Nva, or the N-methylated variant of Ala, Acc, Act, Aib, Apc, or Gly, or of said D- or L- amino acid, or is deleted;

A²³ is Acc, Act, or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Trp, Ser, Thr, Ala, Abu, and Val, or the N-methylated variant of Acc, Act, or Aib, or of said D- or L- amino acid, or is deleted;

A²⁴ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Leu, Ile, Nle, Tle, hLeu, Cha, Val, Ala, Nva, Abu, Trp, and Phe, or the N-methylated variant of Acc or Aib, or of said D- or L- amino acid, or is deleted;

A²⁵ is Acc, Aib, or ~~Aib~~Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Arg, hArg, Lys, Orn, Dab, Dap, Aib, and HN-CH((CH₂)_n-N(R⁴R⁵))-C(O), or the N-methylated variant of Acc, Aib, or ~~Aib~~Apc, or of said D- or L- amino acid, or is deleted;

A²⁶ is Acc, Aib, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of His, 2Pal, 3Pal, 4Pal, Taz, 2Thi, 3Thi, 2Fua, HN-CH((CH₂)_n-N(R⁴R⁵))-C(O), and (X¹,X²,X³,X⁴,X⁵-)Phe, or the N-methylated variant of Acc, Aib, or Apc, or of said D- or L- amino acid, or is deleted;

A²⁷ is Acc or Aic, or a D- or L- amino acid selected from the list of amino acids consisting of Tyr, Phe, hPhe, 2Thi, 3Thi, Taz, 2Fua, Trp, 2Nal, 1Nal, Cha, 2Pal, 3Pal, 4Pal, and (X¹,X²,X³,X⁴,X⁵)Phe, or the N-methylated variant of Acc or Aic or of said D- or L- amino acid;

A²⁸ is Acc or Aib, a D- or L- amino acid selected from the list of amino acids consisting of Leu, Ile, Nle, Tle, hLeu, Trp, Cha, Val, Ala, Nva, Abu, and Phe, or the N-methylated variant of Acc or Aib, or of said D- or L- amino acid;

A²⁹ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Asn, Gln, Glu, Asp, and Trp, or the N-methylated variant of Acc or Aib, or of said D- or L- amino acid;

A³⁰ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Leu, Ile, Nle, Tle, hLeu, Trp, Cha, Val, Ala, Nva, Abu, and Phe or the N-methylated variant of Acc or Aib, or of said D- or L- amino acid;

A³¹ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Val, Leu, Ile, Nle, Tle, hLeu, Cha, Ala, Nva, Abu, Trp, and Phe, or the N-methylated variant of Acc or Aib, or of said D- or L- amino acid;

A³² is Acc, Act, or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Thr, Ser, Ala, Abu, Trp, DTrp, and Val, or the N-methylated variant of Acc, Act, or Aib, or of said D- or L- amino acid;

A³³ is Acc, Aib, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Arg, hArg, Lys, Orn, Dab, Dap, and HN-CH((CH₂)_n-N(R⁴R⁵))-C(O), or the N-methylated variant of Acc, Aib, or Apc, or of said D- or L- amino acid;

A³⁴ is Acc, Aib, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Gln, Asn, Glu, Asp, or the N-methylated variant of Acc, Aib, or Apc, or of said D- or L- amino acid;

A³⁵ is Acc, Aib, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Arg, hArg, Lys, Orn, Dab, Dap, and HN-CH((CH₂)_n-N(R⁴R⁵))-C(O), or the N-methylated variant of Acc, Aib, or Apc, or of said D- or L- amino acid;

A³⁶ is Acc, Aic or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Tyr, Phe, hPhe, 2Thi, 3Thi, Taz, 2Fua, Trp, 2Nal, 1Nal, Cha, 2Pal, 3Pal, 4Pal, and (X¹,X²,X³,X⁴,X⁵)Phe, or the N-methylated variant of Acc, Aic, or Apc, or of said D- or L- amino acid;

R¹ is OH or NH₂, (C₁-C₃₀)alkoxy, or NH-X⁶-CH₂-Z⁰, wherein X⁶ is a (C₁-C₁₂)hydrocarbon moiety, and Z⁰ is -H, -OH, -CO₂H or -C(O)NH₂;

R² and R³ each is, independently for each occurrence, selected from the group consisting of -H, (C₁-C₃₀)alkyl, (C₁-C₃₀)heteroalkyl, (C₁-C₃₀)acyl, (C₂-C₃₀)alkenyl, (C₂-C₃₀)alkynyl, aryl(C₁-C₃₀)alkyl, aryl(C₁-C₃₀)acyl, substituted (C₁-C₃₀)alkyl, substituted (C₁-C₃₀)heteroalkyl, substituted (C₂-C₃₀)acyl, substituted (C₂-C₃₀)alkenyl, substituted (C₂-C₃₀)alkynyl, substituted aryl(C₁-C₃₀)alkyl, and substituted aryl(C₁-C₃₀)acyl,

provided that when R² is (C₁-C₃₀)acyl, aryl(C₁-C₃₀)acyl, substituted (C₂-C₃₀)acyl, or substituted aryl(C₁-C₃₀)acyl, R³ is -H, (C₁-C₃₀)alkyl, (C₁-C₃₀)heteroalkyl, (C₂-C₃₀)alkenyl, (C₂-C₃₀)alkynyl, aryl(C₁-C₃₀)alkyl, substituted (C₁-C₃₀)alkyl, substituted (C₁-C₃₀)heteroalkyl, substituted (C₂-C₃₀)alkenyl, substituted (C₂-C₃₀)alkynyl, or substituted aryl(C₁-C₃₀)alkyl;

R⁴ and R⁵ each is, independently for each occurrence, selected from the group consisting of -H, (C₁-C₄₀)alkyl, (C₂-C₄₀)acyl, (C₁-C₃₀)alkylsulfonyl, and -C(NH)NH₂,

provided that when R⁴ is (C₁-C₄₀)acyl, (C₁-C₃₀)alkylsulfonyl, or -C(NH)NH₂, then R⁵ is -H or (C₁-C₄₀)alkyl;

n is, independently for each occurrence, 1, 2, 3, 4 or 5; and

X^1 , X^2 , X^3 , X^4 , and X^5 each is, independently for each occurrence, selected from the group consisting of -H, -F, -Cl, -Br, -I, (C₁-C₁₀)alkyl, substituted (C₁-C₁₀)alkyl, aryl, substituted aryl, -OH, -NH₂, -NO₂, and -CN;

provided that:

(a) said peptide comprises at least one amino acid selected from the group consisting of:

(i) Acc at A³, A⁶, A⁷, A⁹, A¹⁰, A¹¹, A¹², A¹⁵, A¹⁶, A¹⁷, A¹⁸, A²⁰, A²¹, A²², A²⁴, A²⁷, A²⁸, A²⁹, A³⁰, A³¹, A³², or A³⁴;

(ii) Act at A³, A⁷, A¹², A¹³, A²², A²³, or A³²;

(iii) Apc at A⁴, A⁷, A¹², A¹⁹, A²², A²⁵, A²⁶, A³³, A³⁴, A³⁵, or A³⁶;

(iv) Aib at A⁶, A⁷, A⁹, A¹⁰, A¹¹, A¹², A¹³, A¹⁵, A¹⁶, A¹⁸, A²², A²⁹, or A³²;

(v) Thz, Dmt, Dhp, Ktp, or Tic at A⁵, A⁸, or A¹⁴;

(vi) (3,4,5-F)Phe or (2,3,4,5,6-F)Phe at A²⁰, A²¹, A²⁶, A²⁷, or A³⁶;

(vii) 2Fua at A²⁰, A²¹, A²⁶, or A²⁷;

(viii) Taz at A²⁰, A²¹, or A²⁶; and

(ix) 2Pal, 3Pal, 4Pal, 2Thi or 3Thi at A²⁶;

(b) if A³ - A²¹ are deleted and (i) A²² is Aib or (ii) A³⁶ is (3,4,5-F)Phe or (2,3,4,5,6-F)Phe, then A²⁷ is not 2Thi, Trp, 2Nal, or (X¹,X²,X³,X⁴,X⁵)Phe, wherein X¹ is *p*-chloro and X², X³, X⁴ and X⁵ each is -H; and

(c) each amino acid A^m of formula (I) may be deleted only if A^{m-1} is deleted, wherein m is an integer ranging in value from 4 - 26, inclusive;

or a pharmaceutically acceptable salt thereof.

2. (original) A compound according to claim 1, wherein:

A³ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Ile, Leu, Nle, Tle, hLeu, Cha, Val, Ala, Nva, and Abu, or is deleted;

A⁴ is Acc, Aib, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Lys, Arg, hArg, Orn, Dab, Dap, and HN-CH((CH₂)_n-N(R⁴R⁵))-C(O), or is deleted;

A⁵ is Inc or a D- or L- amino acid selected from the list of amino acids consisting of Pro, Thz, Dmt, Dhp, Ktp, 4Hyp, 3Hyp, Pip, Tic, and Oic, or is deleted;

A⁶ is Acc or a D- or L- amino acid selected from the list of amino acids consisting of Glu, Asp, Gln, Asn, Lys, Arg, Orn, Dab, Dap, and hArg, or is deleted;

A⁷ is Acc, Act, Aib, Apc, or Gly, or a D- or L- amino acid selected from the list of amino acids consisting of Ala, Abu, Val, and Nva, or is deleted;

A⁸ is Inc or a D- or L- amino acid selected from the list of amino acids consisting of Pro, Thz, Dmt, Dhp, Ktp, 4Hyp, 3Hyp, Pip, Tic, and Oic, or is deleted;

A⁹ is Acc, Aib, or Gly or D- or L- Ala, or is deleted;

A¹⁰ is Acc or a D- or L- amino acid selected from the list of amino acids consisting of Glu, Asp, Gln, and Asn, or is deleted;

A¹¹ is Acc or a D- or L- amino acid selected from the list of amino acids consisting of Asp, Glu, Gln, and Asn, or is deleted;

A¹² is Acc, Act, Aib, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Ala, Gly, Abu, Val, and Nva, or is deleted;

A¹³ is Acc, Act, or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Ser, Thr, Ala, Abu, and Val, or is deleted;

A¹⁴ is Inc or a D- or L- amino acid selected from the list of amino acids consisting of Pro, Thz, Dmt, Dhp, Ktp, 4Hyp, 3Hyp, Pip, Tic, and Oic, or is deleted; °

A¹⁵ is Acc or a D- or L- amino acid selected from the list of amino acids consisting of Glu, Asp, Gln, and Asn, or is deleted;

A¹⁶ is Acc or a D- or L- amino acid selected from the list of amino acids consisting of Glu, Asp, Gln, and Asn, or is deleted;

A¹⁷ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Leu, Ile, Nle, Tle, hLeu, Cha, Val, Ala, Nva, Abu, and Phe, or is deleted;

A¹⁸ is Aib or Acc, or a D- or L- amino acid selected from the list of amino acids consisting of Asn, Gln, Glu, and Asp, or is deleted;

A¹⁹ is Acc, Aib, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Arg, hArg, Lys, Orn, Dab, Dap, and HN-CH((CH₂)_n-N(R⁴R⁵))-C(O), or is deleted;

A²⁰ is Acc or Aic, a D- or L- amino acid selected from the list of amino acids consisting of Tyr, Phe, hPhe, 2Thi, 3Thi, Taz, 2Fua, Trp, 2Nal, 1Nal, Cha, 2Pal, 3Pal, 4Pal, and (X¹,X²,X³,X⁴,X⁵)Phe, or is deleted;

A²¹ is Acc or Aic, or a D- or L- amino acid selected from the list of amino acids consisting of Tyr, Phe, hPhe, 2Thi, 3Thi, Taz, 2Fua, Trp, 2Nal, 1Nal, Cha, 2Pal, 3Pal, 4Pal, and (X¹,X²,X³,X⁴,X⁵)Phe, or is deleted;

A²² is Acc, Act, Aib, Apc, or Gly, or a D- or L- amino acid selected from the list of amino acids consisting of Ala, Abu, Val, and Nva, or is deleted;

A²³ is Acc, Act, or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Ser, Thr, Ala, Abu, and Val, or is deleted;

A²⁴ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Leu, Ile, Nle, Tle, hLeu, Cha, Val, Ala, Nva, Abu, Trp, and Phe, or is deleted;

A²⁵ is Acc, Aib, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Arg, hArg, Lys, Orn, Dab, Dap, and HN-CH((CH₂)_n-N(R⁴R⁵))-C(O), or is deleted

A²⁶ is Acc, Aib, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of His, 2Pal, 3Pal, 4Pal, Taz, 2Thi, 3Thi, 2Fua, HN-CH((CH₂)_n-N(R⁴R⁵))-C(O), and (X¹,X²,X³,X⁴,X⁵)-Phe, or is deleted;

A²⁷ is Acc or Aic, or a D- or L- amino acid selected from the list of amino acids consisting of Tyr, Phe, hPhe, 2Thi, 3Thi, Taz, 2Fua, Trp, 2Nal, 1Nal, Cha, 2Pal, 3Pal, 4Pal, and (X¹,X²,X³,X⁴,X⁵)-Phe;

A²⁸ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Leu, Ile, Nle, Tle, hLeu, Trp, Cha, Val, Ala, Nva, Abu, and Phe;

A²⁹ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Asn, Gln, Glu, Asp, and Trp;

A³⁰ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Leu, Ile, Nle, Tle, hLeu, Trp, Cha, Val, Ala, Nva, Abu, and Phe;

A³¹ is Acc or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Val, Leu, Ile, Nle, Tle, hLeu, Cha, Ala, Nva, Abu, Trp, and Phe;

A³² is Acc, Act, or Aib, or a D- or L- amino acid selected from the list of amino acids consisting of Thr, Ser, Ala, Abu, Trp, and Val;

A³³ is Acc, Aib, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Arg, hArg, Lys, Orn, Dab, Dap, and HN-CH((CH₂)_n-N(R⁴R⁵))-C(O);

A³⁴ is Acc, Aib, Apc, or Glu, or a D- or L- amino acid selected from the list of amino acids consisting of Gln, Asn, and Asp;

A³⁵ is Acc, Aib, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Arg, hArg, Lys, Orn, Dab, Dap, and HN-CH((CH₂)_n-N(R⁴R⁵))-C(O); and

A³⁶ is Acc, Aic, or Apc, or a D- or L- amino acid selected from the list of amino acids consisting of Tyr, Phe, hPhe, 2Thi, 3Thi, Taz, 2Fua, Trp, 2Nal, 1Nal, Cha, 2Pal, 3Pal, 4Pal, and (X¹,X²,X³,X⁴,X⁵)-Phe;

or a pharmaceutically acceptable salt thereof.

3. (original) A compound according to claim 2, wherein:

A³ is Ile, Leu, Nle, Tle, hLeu, Cha, Val, Ala, Nva, Abu, Acc, or Aib, or is deleted;

A⁴ is Lys, Arg, hArg, Orn, Dab, Dap, Apc, Aib, Acc, or HN-CH((CH₂)_n-N(R⁴R⁵))-C(O), or is deleted;

A⁵ is Pro, Thz, Dmt, Dhp, Ktp, 4Hyp, 3Hyp, Pip, Tic, Oic, or Inc, or is deleted;

A⁶ is Glu, Asp, Gln, Asn, Lys, Arg, Orn, Dab, Dap, hArg, or Acc, or is deleted;

A⁷ is Ala, Aib, Gly, Abu, Val, Nva, Apc, Act, or Acc, or is deleted;

A⁸ is Pro, Thz, Dmt, Dhp, Ktp, 4Hyp, 3Hyp, Pip, Tic, Oic, or Inc, or is deleted;

A⁹ is Gly, Ala, Aib, or Acc, or is deleted;

A¹⁰ is Glu, Asp, Gln, Asn, or Acc, or is deleted;

A¹¹ is Asp, Glu, Gln, Asn, or Acc, or is deleted;

A¹² is Ala, Aib, Gly, Abu, Val, Nva, Apc, Act, or Acc, or is deleted;

A¹³ is Ser, Thr, Aib, Act, Ala, Acc, Abu, or Val, or is deleted;

A¹⁴ is Pro, Thz, Dmt, Dhp, Ktp, 4Hyp, 3Hyp, Pip, Tic, Oic, or Inc, or is deleted;

A¹⁵ is Glu, Asp, Gln, Asn, or Acc, or is deleted;

A¹⁶ is Glu, Asp, Gln, Asn, or Acc, or is deleted;

A¹⁷ is Leu, Ile, Nle, Tle, hLeu, Cha, Val, Ala, Nva, Abu, Acc, Aib, or Phe, or is deleted;

A¹⁸ is Asn, Gln, Glu, Asp, Aib, or Acc, or is deleted;

A¹⁹ is Arg, hArg, Lys, Orn, Dab, Dap, Apc, Aib, Acc, or HN-CH((CH₂)_n-N(R⁴R⁵))-C(O), or is deleted;

A²⁰ is Tyr, Phe, hPhe, 2Thi, 3Thi, Taz, 2Fua, Trp, 2Nal, 1Nal, Cha, 2Pal, 3Pal, 4Pal, (X¹,X²,X³,X⁴,X⁵)Phe, Acc, or Aic, or is deleted;

A²¹ is Tyr, Phe, hPhe, 2Thi, 3Thi, Taz, 2Fua, Trp, 2Nal, 1Nal, Cha, 2Pal, 3Pal, 4Pal, (X¹,X²,X³,X⁴,X⁵)Phe, Acc, or Aic, or is deleted;

A²² is Ala, Aib, Gly, Abu, Val, Nva, Apc, Act, Acc, or N-Me-Ala, or is deleted;

A²³ is Ser, Thr, Aib, Act, Ala, Acc, Abu, Val, or DTrp, or is deleted;

A²⁴ is Leu, Ile, Nle, Tle, hLeu, Cha, Val, Ala, Nva, Abu, Acc, Aib, Trp, or Phe, or is deleted;

A²⁵ is Arg, hArg, Lys, Orn, Dab, Dap, Apc, Aib, HN-CH((CH₂)_n-N(R⁴R⁵))-C(O), or Acc, or is deleted;

A²⁶ is His, 2Pal, D2Pal, 3Pal, 4Pal, Taz, 2Thi, 3Thi, 2Fua, Apc, Aib, Acc, HN-CH((CH₂)_n-N(R⁴R⁵))-C(O), or (X¹,X²,X³,X⁴,X⁵)Phe, or is deleted;

A²⁷ is Tyr, Phe, hPhe, 2Thi, 3Thi, Taz, 2Fua, Trp, 2Nal, 1Nal, Cha, 2Pal, 3Pal, 4Pal, (X¹,X²,X³,X⁴,X⁵)Phe, Acc, or Aic;

A²⁸ is Leu, Ile, Nle, Tle, hLeu, Trp, Cha, Val, Ala, Nva, Abu, Acc, Aib, or Phe;

A²⁹ is Asn, Gln, Glu, Asp, Acc, Trp, or Aib;

A³⁰ is Leu, Ile, Nle, Tle, hLeu, Trp, Cha, Val, Ala, Nva, Abu, Acc, Aib, or Phe;

A^{31} is Val, Leu, Ile, Nle, Tle, hLeu, Cha, Ala, Nva, Abu, Acc, Aib, Trp, or Phe;
 A^{32} is Thr, Ser, Aib, Act, Ala, Acc, Abu, Trp, DTrp, or Val;
 A^{33} is Arg, hArg, Lys, Orn, Dab, Dap, Apc, Aib, $\text{HN-CH}((\text{CH}_2)_n\text{-N}(\text{R}^4\text{R}^5))\text{-C}(\text{O})$, or Acc;
 A^{34} is Gln, Asn, Glu, Asp, Acc, Aib, or Apc;
 A^{35} is Arg, hArg, Lys, Orn, Dab, Dap, Apc, Aib, $\text{HN-CH}((\text{CH}_2)_n\text{-N}(\text{R}^4\text{R}^5))\text{-C}(\text{O})$, or Acc; and
 A^{36} is Tyr, Phe, hPhe, 2Thi, 3Thi, Taz, 2Fua, Trp, 2Nal, 1Nal, Cha, 2Pal, 3Pal, 4Pal, $(\text{X}^1, \text{X}^2, \text{X}^3, \text{X}^4, \text{X}^5)\text{Phe}$, Acc, Aic, or Apc; or
 a pharmaceutically acceptable salt thereof.

4. (original) A compound according to claim 3, wherein:

A^3 is Ile, Leu, Nle, Val, Acc, or Aib, or is deleted;
 A^4 is Lys, Arg, hArg, Orn, or Apc, or is deleted;
 A^5 is Pro, Thz, Dmt, 4Hyp, or 3Hyp, or is deleted;
 A^6 is Glu, Asp, Gln, or Acc, or is deleted;
 A^7 is Ala, Aib, Abu, Act, or Acc, or is deleted;
 A^8 is Pro, Thz, Dmt, 4Hyp, or 3Hyp, or is deleted;
 A^9 is Gly, Aib, or Acc, or is deleted;
 A^{10} is Glu, Asp, Gln, or Acc or is deleted;
 A^{11} is Asp, Glu, Asn, or Acc or is deleted;
 A^{12} is Ala, Aib, Act, or Acc, or is deleted;
 A^{13} is Ser, Thr, Aib, Act, or Acc, or is deleted;
 A^{14} is Pro, Thz, Dmt, 4Hyp, or 3Hyp, or is deleted;
 A^{15} is Glu, Asp, Gln, or Acc, or is deleted;
 A^{16} is Glu, Asp, Gln, or Acc or is deleted;
 A^{17} is Leu, Ile, Nle, Val, Acc, or Aib, or is deleted;
 A^{18} is Asn, Gln, Asp, Aib, or Acc or is deleted;
 A^{19} is Arg, hArg, Lys, or Apc, or is deleted;
 A^{20} is Tyr, Phe, 2Pal, 3Pal, 4Pal, $(\text{X}^1, \text{X}^2, \text{X}^3, \text{X}^4, \text{X}^5)\text{Phe}$, or Acc, or is deleted;
 A^{21} is Tyr, Phe, 2Pal, 3Pal, 4Pal, $(\text{X}^1, \text{X}^2, \text{X}^3, \text{X}^4, \text{X}^5)\text{Phe}$, or Acc, or is deleted;
 A^{22} is Ala, Aib, Abu, or Acc, or is deleted;
 A^{23} is Ser, Thr, Aib, Act, or Ala, or is deleted;
 A^{24} is Leu, Ile, Nle, Val, Acc, or Aib, or is deleted;
 A^{25} is Arg, hArg, Lys, or Apc, or is deleted;

A²⁶ is His, 2Pal, D2Pal, 3Pal, 4Pal, Taz, 2Thi, 3Thi, Apc, or (X¹,X²,X³,X⁴,X⁵-)Phe, or is deleted;

A²⁷ is Tyr, Phe, 2Pal, 3Pal, 4Pal, (X¹,X²,X³,X⁴,X⁵)Phe or Acc;

A²⁸ is Leu, Ile, Nle, Val, Acc or Aib;

A²⁹ is Asn, Gln, Asp, Acc or Aib;

A³⁰ is Leu, Ile, Nle, Val, Acc or Aib;

A³¹ is Val, Leu, Ile, Ala, Acc or Aib;

A³² is Thr, Ser, Aib, Act or Acc;

A³³ is Arg, hArg, Lys or Apc;

A³⁴ is Gln, Asn, Glu, Aib or Apc;

A³⁵ is Arg, hArg, Lys or Apc; and

A³⁶ is Tyr, Phe, 2Pal, 3Pal, 4Pal, (X¹,X²,X³,X⁴,X⁵)Phe or Apc;

or a pharmaceutically acceptable salt thereof.

5. (original) A compound according to claim 4, wherein:

A³ is Ile or Acc, or is deleted;

A⁴ is Lys or Apc, or is deleted;

A⁵ is Pro or is deleted;

A⁶ is Glu or Acc, or is deleted;

A⁷ is Ala, Act, or Acc, or is deleted;

A⁸ is Pro or is deleted;

A⁹ is Gly or Acc, or is deleted;

A¹⁰ is Glu or Acc, or is deleted;

A¹¹ is Asp or Acc, or is deleted;

A¹² is Ala, Act, or Acc, or is deleted;

A¹³ is Ser, Act, or Acc, or is deleted;

A¹⁴ is Pro or is deleted;

A¹⁵ is Glu or Acc, or is deleted;

A¹⁶ is Glu or Acc, or is deleted;

A¹⁷ is Leu or Acc, or is deleted;

A¹⁸ is Asn or Acc, or is deleted;

A¹⁹ is Arg or Apc, or is deleted;

A²⁰ is Tyr, (X¹,X²,X³,X⁴,X⁵)Phe, or Acc, or is deleted;

A²¹ is Tyr, (X¹,X²,X³,X⁴,X⁵)Phe, or Acc, or is deleted;

A^{22} is Ala, Aib, or Acc, or is deleted;
 A^{23} is Ser or Act, or is deleted;
 A^{24} is Leu or Acc, or is deleted;
 A^{25} is Arg or Apc, or is deleted;
 A^{26} is His, 2Pal, D2Pal, 3Pal, 4Pal, Taz, Apc, or $(X^1, X^2, X^3, X^4, X^5)$ Phe, or is deleted;
 A^{27} is Tyr, $(X^1, X^2, X^3, X^4, X^5)$ Phe, or Acc;
 A^{28} is Leu, or Acc;
 A^{29} is Asn or Acc;
 A^{30} is Leu or Acc;
 A^{31} is Val, Leu or Acc;
 A^{32} is Thr, Act, or Acc;
 A^{33} is Arg or Apc;
 A^{34} is Gln or Apc;
 A^{35} is Arg or Apc; and
 A^{36} is Tyr, $(X^1, X^2, X^3, X^4, X^5)$ Phe, or Apc;

or a pharmaceutically acceptable salt thereof.

6. (original) A compound according to claim 5, wherein:

Acc is, independently for each occurrence, A5c or A6c; and

$(X^1, X^2, X^3, X^4, X^5)$ Phe is, independently for each occurrence, (3,4,5-F)Phe or (2,3,4,5,6-F)Phe;

or a pharmaceutically acceptable salt thereof.

7. (original) A compound according to claim 6, wherein:

A^3 is Ile or is deleted;

A^4 is Lys or is deleted;

A^6 is Glu or is deleted;

A^7 is Ala or is deleted;

A^9 is Gly or is deleted;

A^{10} is Glu or is deleted;

A^{11} is Asp or is deleted;

A^{12} is Ala or is deleted;

A^{13} is Ser or is deleted;

A^{14} is Pro or is deleted;

A^{15} is Glu or is deleted;

A^{16} is Glu or is deleted;
 A^{17} is Leu or is deleted;
 A^{18} is Asn or is deleted;
 A^{19} is Arg or is deleted;
 A^{20} is Tyr or is deleted;
 A^{21} is Tyr or is deleted;
 A^{22} is Ala, Aib, or A5c, or is deleted;
 A^{23} is Ser or is deleted;
 A^{24} is Leu or A6c;
 A^{25} is Arg;
 A^{26} is His, 2Pal, D2Pal, 3Pal, 4Pal, or Taz;
 A^{27} is Tyr or (3,4,5-F)Phe;
 A^{28} is Leu, or A6c;
 A^{29} is Asn;
 A^{30} is Leu or A6c;
 A^{31} is Val, Leu, A5c or A6c;
 A^{32} is Thr;
 A^{33} is Arg;
 A^{34} is Gln; and
 A^{36} is Tyr;

or a pharmaceutically acceptable salt thereof.

8. (original) A compound according to claim 6, wherein said compound is according to the formula:

$((2,3,4,5,6\text{-F})\text{Phe}^{20})\text{hPYY}(3\text{-}36)\text{NH}_2$;	(SEQ ID NO. 31)
$((2,3,4,5,6\text{-F})\text{Phe}^{21})\text{hPYY}(3\text{-}36)\text{NH}_2$;	(SEQ ID NO. 32)
$\text{Ac}-((2,3,4,5,6\text{-F})\text{Phe}^{26})\text{hPYY}(22\text{-}36)\text{NH}_2$;	(SEQ ID NO. 33)
$\text{Ac}-((2,3,4,5,6\text{-F})\text{Phe}^{26})\text{hPYY}(24\text{-}36)\text{NH}_2$;	(SEQ ID NO. 34)
$((2,3,4,5,6\text{-F})\text{Phe}^{26})\text{hPYY}(3\text{-}36)\text{NH}_2$;	(SEQ ID NO. 35)
$\text{Ac}-((2,3,4,5,6\text{-F})\text{Phe}^{27})\text{hPYY}(22\text{-}36)\text{NH}_2$;	(SEQ ID NO. 36)
$\text{Ac}-((2,3,4,5,6\text{-F})\text{Phe}^{27})\text{hPYY}(24\text{-}36)\text{NH}_2$;	(SEQ ID NO. 37)
$((2,3,4,5,6\text{-F})\text{Phe}^{27})\text{hPYY}(3\text{-}36)\text{NH}_2$;	(SEQ ID NO. 38)
$\text{Ac}-((2,3,4,5,6\text{-F})\text{Phe}^{36})\text{hPYY}(22\text{-}36)\text{NH}_2$;	(SEQ ID NO. 39)
$\text{Ac}-((2,3,4,5,6\text{-F})\text{Phe}^{36})\text{hPYY}(24\text{-}36)\text{NH}_2$;	(SEQ ID NO. 40)
$((2,3,4,5,6\text{-F})\text{Phe}^{36})\text{hPYY}(3\text{-}36)\text{NH}_2$;	(SEQ ID NO. 41)

((3,4,5-F)Phe ²⁰)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 42)
((3,4,5-F)Phe ²¹)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 43)
Ac-((3,4,5-F)Phe ²⁶)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 44)
Ac-((3,4,5-F)Phe ²⁶)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 45)
((3,4,5-F)Phe ²⁶)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 46)
Ac-((3,4,5-F)Phe ²⁷)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 15)
Ac-((3,4,5-F)Phe ²⁷)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 47)
((3,4,5-F)Phe ²⁷)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 12)
Ac-((3,4,5-F)Phe ³⁶)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 48)
Ac-((3,4,5-F)Phe ³⁶)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 49)
((3,4,5-F)Phe ³⁶)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 50)
Ac-(D2Pal ²⁶)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 26)
Ac-(2Pal ²⁶)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 27)
Ac-(2Pal ²⁶ , Leu ³¹)hPPY(24-36)NH ₂ ;	(SEQ ID NO. 18)
Ac-(3Pal ²⁶)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 14)
(3Pal ²⁶)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 5)
Ac-(3Pal ²⁶ , Leu ³¹)hPPY(24-36)NH ₂ ;	(SEQ ID NO. 16)
Ac-(4Pal ²⁶)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 13)
Ac-(4Pal ²⁶ , Leu ³¹)hPPY(24-36)NH ₂ ;	(SEQ ID NO. 17)
Ac-(A5c ²²)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 4)
Ac-(A5c ³¹)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 24)
Ac-(A5c ³¹)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 51)
(A5c ³¹)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 3)
(A6c ¹⁰)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 52)
(A6c ¹¹)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 53)
(A6c ¹²)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 54)
(A6c ¹³)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 55)
(A6c ¹⁵)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 56)
(A6c ¹⁶)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 57)
(A6c ¹⁷)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 58)
(A6c ¹⁸)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 59)
(A6c ²⁰)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 60)
(A6c ²¹)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 61)
Ac-(A6c ²²)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 62)

(A6c ²²)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 63)
Ac-(A6c ²⁴)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 25)
Ac-(A6c ²⁴)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 64)
(A6C ²⁴)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 10)
Ac-(A6c ²⁴ , Leu ³¹)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 28)
Ac-(A6c ²⁷)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 65)
Ac-(A6c ²⁷)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 66)
(A6c ²⁷)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 67)
Ac-(A6c ²⁸)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 23)
Ac-(A6c ²⁸)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 68)
(A6c ²⁸)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 8)
Ac-(A6c ²⁸ , Leu ³¹)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 29)
Ac-(A6c ²⁹)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 69)
Ac-(A6c ²⁹)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 70)
(A6c ²⁹)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 71)
(A6c ³)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 72)
Ac-(A6c ³⁰)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 22)
Ac-(A6c ³⁰)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 73)
(A6c ³⁰)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 9)
Ac-(A6c ³¹)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 21)
Ac-(A6c ³¹)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 30)
(A6c ³¹)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 74)
Ac-(A6c ³²)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 75)
Ac-(A6c ³²)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 76)
(A6c ³²)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 77)
(A6c ⁶)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 78)
(A6c ⁷)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 79)
(A6c ⁹)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 80)
(Act ¹²)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 81)
(Act ¹³)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 82)
Ac-(Act ²³)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 83)
(Act ²³)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 84)
Ac-(Act ³²)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 85)
Ac-(Act ³²)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 86)

(Act ³²)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 87)
(Act ⁷)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 88)
Ac-(Aib ²²)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 89)
(Aib ²²)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 11)
(Apc ¹⁹)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 90)
Ac-(Apc ²⁵)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 91)
Ac-(Apc ²⁵)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 92)
(Apc ²⁵)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 93)
Ac-(Apc ²⁶)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 94)
Ac-(Apc ²⁶)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 95)
(Apc ²⁶)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 96)
Ac-(Apc ³³)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 97)
Ac-(Apc ³³)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 98)
(Apc ³³)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 99)
Ac-(Apc ³⁴)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 100)
Ac-(Apc ³⁴)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 101)
(Apc ³⁴)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 102)
Ac-(Apc ³⁵)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 103)
Ac-(Apc ³⁵)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 104)
(Apc ³⁵)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 7)
Ac-(Apc ³⁶)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 105)
Ac-(Apc ³⁶)hPYY(24-36)NH ₂ ;	(SEQ ID NO. 106)
(Apc ³⁶)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 107)
(Apc ⁴)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 108)
(Taz ²⁶)hPYY(3-36)NH ₂ ;	(SEQ ID NO. 6)
Ac-(Taz ²⁶)hPYY(22-36)NH ₂ ;	(SEQ ID NO. 20)
Ac-(Taz ²⁶ , Leu ³¹)hPPY(24-36)NH ₂ ;	(SEQ ID NO. 19)

or a pharmaceutically acceptable salt thereof.

9. (currently amended) A compound according to claim 8, wherein said compound is according to the formula:

[A5C ³¹]hPYY(3-36)NH ₂	(SEQ ID NO. 3)
Ac-[A5C ²²]hPYY(22-36)NH ₂	(SEQ ID NO. 4)
[3Pal ²⁶]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 5)

[Taz ²⁶]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 6)
[Apc ³⁵]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 7)
[A6C ²⁸]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 8)
[A6C ³⁰]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 9)
[A6C ²⁴]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 10)
[Aib ²²]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 11)
[((3,4,5-F)Phe) ²⁷]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 12)
Ac-[4Pal ²⁶]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 13)
Ac-[3Pal ²⁶]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 14)
Ac-(((3,4,5-F)Phe) ²⁷]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 15)
Ac-(3Pal ²⁶ , Leu ³¹)hPPY(24-36)NH ₂ ;	(SEQ ID NO. 16)
Ac-(4Pal ²⁶ , Leu ³¹)hPPY(24-36)NH ₂ ;	(SEQ ID NO. 17)
Ac-(2Pal ²⁶ , Leu ³¹)hPPY(24-36)NH ₂ ;	(SEQ ID NO. 18)
Ac-(Taz ²⁶ , Leu ³¹)hPPY(24-36)NH ₂ ;	(SEQ ID NO. 19)
Ac-[Taz ²⁶]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 20)
Ac-[A6c ³¹]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 21)
Ac-[A6c ³⁰]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 22)
Ac-[A6c ²⁸]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 23)
Ac-[A5c ³¹]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 24)
Ac-[A6C ²⁴]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 25)
Ac-[D2Pal ²⁶]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 26)
Ac-[2Pal ²⁶]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 27)
Ac-[A6C ²⁴ , Leu ³¹]hPYY(24-36)NH ₂ ;	(SEQ ID NO. 28)
Ac-[A6C ²⁸ , Leu ³¹]hPYY(24-36)NH ₂ ;	(SEQ ID NO. 29)
Ac-[A6C ³¹]hPYY(24-36)NH ₂ ;	(SEQ ID NO. 30)
<u>Ac-(A6c²⁴)hPYY(24-36)NH₂;</u>	<u>(SEQ ID NO. 64)</u>

or a pharmaceutically acceptable salt thereof.

10. (original) A compound according to claim 9, wherein said compound is:

[A5C ³¹]hPYY(3-36)NH ₂	(SEQ ID NO. 3)
[3Pal ²⁶]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 5)
[Taz ²⁶]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 6)
[A6C ²⁸]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 8)
[A6C ²⁴]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 10)

[Aib ²²]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 11)
[((3,4,5-F)Phe) ²⁷]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 12)
Ac-[4Pal ²⁶]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 13)
Ac-[3Pal ²⁶]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 14)
Ac-(3Pal ²⁶ , Leu ³¹)hPPY(24-36)NH ₂ ;	(SEQ ID NO. 16)
Ac-(4Pal ²⁶ , Leu ³¹)hPPY(24-36)NH ₂ ;	(SEQ ID NO. 17)

or a pharmaceutically acceptable salt thereof.

11. (original) A compound according to claim 9, wherein said compound is:

[A5C ³¹]hPYY(3-36)NH ₂	(SEQ ID NO. 3)
[3Pal ²⁶]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 5)
[Taz ²⁶]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 6)
[Apc ³⁵]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 7)
[A6C ²⁸]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 8)
[A6C ²⁴]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 10)
[Aib ²²]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 11)
Ac-[4Pal ²⁶]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 13)
Ac-[3Pal ²⁶]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 14)
Ac-(3Pal ²⁶ , Leu ³¹)hPPY(24-36)NH ₂ ;	(SEQ ID NO. 16)
Ac-(4Pal ²⁶ , Leu ³¹)hPPY(24-36)NH ₂ ;	(SEQ ID NO. 17)

or a pharmaceutically acceptable salt thereof.

12. (currently amended) A compound according to claim 9, wherein said compound is:

[A5C ³¹]hPYY(3-36)NH ₂	(SEQ ID NO. 3)
[3Pal ²⁶]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 5)
[A6C ²⁸]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 8)
[A6C ²⁴]hPYY(3-36)NH ₂ ;	(SEQ ID NO. 10)
Ac-[4Pal ²⁶]hPYY(22-36)NH ₂ ;	(SEQ ID NO. 13)
<u>Ac-(A6c²⁴)hPYY(24-36)NH₂;</u>	<u>(SEQ ID NO. 64)</u>

or a pharmaceutically acceptable salt thereof.

13. (currently amended) A pharmaceutical composition comprising a compound according to claim 1 ~~any one of claims 1—12~~, or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier.

14. (currently amended) A method of decreasing excess intestinal water and electrolyte secretion in a mammal in need thereof, said method comprising administering to said mammal an effective amount of a compound according to claim 1 ~~any one of claims 1—12~~, or a pharmaceutically acceptable salt thereof.
15. (currently amended) A method of regulating cell proliferation in a mammal in need thereof, said method comprising administering to said mammal an effective amount of a compound according to claim 1 ~~any one of claims 1—12~~, or a pharmaceutically acceptable salt thereof.
16. (original) A method of claim 15, wherein said cell is a gastrointestinal cell.
17. (original) A method of claim 15, wherein said cell is an epithelial cell.
18. (currently amended) A method of augmenting nutrient transport in a mammal in need thereof, said method comprising administering to said mammal an effective amount of a compound according to claim 1 ~~any one of claims 1—12~~, or a pharmaceutically acceptable salt thereof.
19. (currently amended) A method of regulating lipolysis in a mammal in need thereof, said method comprising administering to said mammal an effective amount of a compound according to claim 1 ~~any one of claims 1—12~~, or a pharmaceutically acceptable salt thereof.
20. (currently amended) A method of regulating blood flow in a mammal in need thereof, said method comprising administering to said mammal an effective amount of a compound according to claim 1 ~~any one of claims 1—12~~, or a pharmaceutically acceptable salt thereof.
21. (currently amended) A method of facilitating weight loss, appetite decrease, weight maintenance, treating obesity, treating diabetes, treating complications of diabetes including retinopathy, or treating cardiovascular disorders in a mammal in need thereof, said method comprising administering to said mammal an effective amount of a compound according to claim 1 ~~any one of claims 1—12~~, or a pharmaceutically acceptable salt thereof.
22. (original) A method according to claim 21, wherein excessive weight is a contributing factor to a disease or condition including hypertension, diabetes, dyslipidemia, cardiovascular disease, gall stones, osteoarthritis and cancers.

23. (original) A method according to claim 22, wherein said facilitation of weight loss reduces the likelihood of such diseases or conditions or where said facilitation of weight loss comprises at least part of a treatment for such diseases or conditions.

24. (currently amended) A method of antagonizing the effects of PYY(3-36) in a mammal in need thereof, said method comprising administering to said mammal an effective amount of a compound according to claim 1~~any one of claims 1—12~~, or a pharmaceutically acceptable salt thereof, wherein said compound is a PYY antagonist.

25. (original) A method according to claim 24, wherein said antagonist effects in said mammal comprise facilitating weight gain, facilitating maintenance in weight, and/or facilitating appetite increase.

26. (original) A method according to claim 25, wherein said facilitating weight gain, facilitating maintenance in weight, and/or facilitating appetite increase is indicated in a mammal having a disease or disorder, or under going a treatment, accompanied by weight loss.

27. (original) A method according to claim 26, wherein said diseases or disorders accompanied by weight loss include anorexia, bulimia, cancer cachexia, AIDS, wasting, cachexia, and wasting in frail elderly.

28. (original) A method according to claim 26, wherein said treatment accompanied by weight loss comprises chemotherapy, radiation therapy, temporary or permanent immobilization, or dialysis.